

Date: 11/13/00 Express Mail Label No. EL552577906 US

Inventors: C. Thomas Caskey, John Shumaker and Andres Metspalu  
5 Attorney's Docket No.: 2875.1001-007

## PARALLEL PRIMER EXTENSION APPROACH TO NUCLEIC ACID SEQUENCE ANALYSIS

### RELATED APPLICATIONS

10 This application is a Continuation of United States Application No. 08/564,100,  
which is the United States National Phase Application of PCT/US94/07086, filed June  
22, 1994 which is a Continuation-in-Part Application of Sweden Application No. SE  
9302152-5, filed on June 22, 1993, the entire teachings of which are incorporated herein  
by reference in their entireties.

### GOVERNMENT FUNDING

15 This invention was made with Government Support under grant number 5-R01-  
DK31 428-11 awarded by the National Institutes of Health. The United States  
Government has certain rights in the invention.

### BACKGROUND OF THE INVENTION

20 Today, there are two predominant methods for DNA sequence determination:  
the chemical degradation method (Maxam and Gilbert, Proc. Natl. Acad. Sci. 74:560-  
564 (1977), and the dideoxy chain termination method (Sanger et al., Proc. Natl. Acad.  
Sci. 74:5463-5467 (1977)). Most automated sequencers are based on the chain  
termination method utilizing fluorescent detection of product formation. There are two  
common variations of these systems: (1) dye-labeled primers to which  
25 deoxynucleotides and dideoxynucleotides are added, and (2) primers to which  
deoxynucleotides and fluorescently labeled dideoxynucleotides are added. In addition,  
the labeled deoxynucleotides can be used in conjunction with unlabeled  
dideoxynucleotides. This method is based upon the ability of an enzyme to add specific

DOCKET = 2875.1001-007

Although both the chemical degradation method and the dideoxy chain termination method are in widespread use, there are many associated disadvantages: for example, the methods require gel-electrophoretic separation. Typically, only 400-800 base pairs can be sequenced from a single clone. As a result, the systems are both time- and labor-intensive. Methods avoiding gel separation have been developed in attempts to increase the sequencing throughput.

These methods are problematic in several respects. For example, the hybridization is dependent upon the sequence composition of the duplex of the oligonucleotide and the polynucleotide of interest, so that GC-rich regions are more stable than AT-rich regions. As a result, false positives and false negatives during hybridization detection are frequently present and complicate sequence determination. Furthermore, the sequence of the polynucleotide is not determined directly, but is inferred from the sequence of the known probe, which increases the possibility for error. A great need remains to develop efficient and accurate methods for nucleic acid sequence determination.

The current invention pertains to methods for analyzing, and particularly for sequencing, a polynucleotide of interest, and an apparatus useful in analyzing a polynucleotide of interest. In one embodiment of the current invention, the nucleotide sequence of a polypeptide of interest is analyzed for the presence of mutations or

alterations. In a second embodiment of the current invention, the nucleotide sequence of a polypeptide of interest, for which the nucleotide sequence was not known previously, is determined. The method comprises detecting single base extension events of a set of specific oligonucleotide primers, such that the label and position of each  
5 separate extension event defines a base in a polynucleotide of interest.

In one method of the current invention, a solid support is provided. An array of a set or several sets of consecutive oligonucleotide primers of a specified size having known sequences is attached at defined locations to the solid support. The oligonucleotide primers differ within each set by one base pair. The oligonucleotide  
10 primers either correspond to at least a part of the nucleotide sequence of one strand of the polynucleotide of interest, if the sequence is known, or represent a set of all possible nucleotide sequences for oligonucleotide primers of the specified size, if the sequence is not known. A polynucleotide of interest, which may be DNA or RNA, or a fragment of the polynucleotide of interest, is annealed to the array of oligonucleotide primers under  
15 hybridization conditions, thereby generating "annealed primers". The annealed primers are subjected to single base extension reaction conditions, under which a nucleic acid polymerase and terminating nucleotides, such as dideoxynucleotides (ddNTPs) corresponding to the four known bases (A, G, T and C), are provided to the annealed primers. The terminating nucleotides can also comprise a terminating string of known  
20 polynucleotides, such as dinucleotides. As a result of the single base extension reaction, extended primers are generated, in which a terminating nucleotide is added to each of the annealed primers. The terminating nucleotides can be provided to the annealed primers either simultaneously or sequentially. The terminating nucleotides are mutually distinguishable; i.e., at least one of the nucleotides is labelled to facilitate detection.  
25 After addition of the terminating nucleotides, the sequence of the polynucleotide of interest is analyzed by "reading" the oligonucleotide array: the identity and location of each terminating nucleotide within the array on the solid support is observed. The label and position of each terminating nucleotide on the solid support directly defines the sequence of the polynucleotide of interest that is being analyzed.

In a second method of the current invention, the polynucleotide of interest is  
30 analyzed for the presence of specific mutations through the use of oligonucleotide primers that are not attached to a solid support. The oligonucleotide primers are tailored to anneal to the polynucleotide of interest at a point immediately preceding the mutation site(s). If more than one mutation site is examined, the oligonucleotide primers are  
35 designed to be mutually distinguishable: in a preferred embodiment, the

oligonucleotide primers have different mobilities during gel electrophoresis. For example, oligonucleotides of different lengths are used. After the oligonucleotide primers are annealed to the polynucleotide of interest, the annealed primers are subjected to single base extension reaction conditions, resulting in extended primers in which terminating nucleotides are added to each of the annealed primers. As in the first method of the current invention, the terminating nucleotides are mutually distinguishable. After addition of the terminating nucleotides, the sequence of the polynucleotide of interest is analyzed by eluting the extended primers, performing gel electrophoresis, and "reading" the gel: the identity and location of each terminating nucleotides on the gel is observed using standard methods, such as with an automated DNA sequencer. The label and position of each terminating nucleotide on the gel directly defines the sequence of the polynucleotide of interest that is being analyzed, and indicates whether a mutation is present.

The apparatus of the current invention comprises a solid support having an array of one or more sets of consecutive oligonucleotide primers with known sequences attached to it at defined locations, each oligonucleotide primer differing within each set by one base pair. The set of oligonucleotide primers either corresponds to at least a part of the nucleotide sequence of one strand of the polynucleotide of interest, if the sequence is known, or represents all possible nucleotide sequences for oligonucleotide primers of the specified size, if the sequence is not known.

The current invention provides both direct information, due to the detection of a specific nucleotide addition, and indirect information, due to the known sequence of the annealed primer to which the specific base addition occurred, for the polynucleotide of interest. The ability to determine nucleic acid sequences is a critical element of understanding gene expression and regulation. In addition, as advances in molecular medicine continue, sequence determination will become a more important element in the diagnosis and treatment of disease.

#### BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 depicts an example of a set of oligonucleotide primers (sense primers, SEQ ID NOS: 2-11, antisense primers, SEQ ID NOS: 12-21), comprising consecutive primers differing by one base pair at the growing end and capable of hybridizing successively along the relevant part(s) or the whole of the polynucleotide of interest (SEQ ID NOS: 1 and 22).

Figure 2 is a schematic illustration of a single strand template bound to a primer which is in turn attached to a solid support.

Figure 3 illustrates a set of consecutive oligonucleotide primers for a part of the polynucleotide of interest following immediately after the primer illustrated in Figure 2.

5 Figure 4 illustrates the single base pair additions to all the primers illustrated in Figure 3, as well as the corresponding additions for the corresponding primers related to the complementary strand of the polynucleotide of interest.

Figure 5 is a graphic depiction of the length of extended primers formed utilizing free oligonucleotide primers annealed to a polynucleotide of interest.

10 Figures 6A, 6B and 6C are graphic depictions of electrophoretograms demonstrating the detection of the presence of a mutation in a polynucleotide of interest.

Figures 7A, 7B and 7C depict the results of a DNA chip-based analysis for a five-base region within the third exon of the HPRT gene.

#### DETAILED DESCRIPTION OF THE INVENTION

15 The current invention pertains to methods for analyzing the nucleotide sequence of a polynucleotide of interest. The method comprises hybridizing all or a fragment of a polynucleotide of interest to oligonucleotide primers, conducting single base extension reactions, and detecting the single base extension events. The method can be used to analyze the sequence of a polypeptide of interest by examining the sequence for the  
20 presence of mutations or alterations in the nucleotide sequence, or by determining the nucleotide sequence of a polypeptide of interest.

As used herein, the term "polynucleotide of interest" refers to the particular polynucleotide for which sequence information is wanted. Representative polynucleotides of interest include oligonucleotides, DNA or DNA fragments, RNA or  
25 RNA fragments, as well as genes or portions of genes. The polynucleotide of interest can be single- or double-stranded. The term "template polynucleotide of interest" is used herein to refer to the strand which is analyzed, if only one strand of a double-stranded polynucleotide is analyzed, or to the strand which is identified as the first strand, if both strands of a double-stranded polynucleotide are analyzed. The term,  
30 "complementary polynucleotide of interest" is used herein to refer to the strand which is not analyzed, if only one strand of a double-stranded polynucleotide is analyzed, or to the strand which is identified as the second strand (i.e., the strand that is complementary to the first (template) strand), if both strands of a double-stranded polynucleotide are analyzed. Either one of the two strands can be analyzed. In a preferred embodiment,

0074475:4430

both strands of a double-stranded polynucleotide of interest are analyzed in order to verify sequence information obtained from the template (first) strand by comparison with the complementary (second) strand. Nevertheless, it is not always necessary to analyze both strands. For example, if the polynucleotide of interest is being analyzed for the presence of a single base mutation, and not for the complete base sequence in the mutation region, it is sufficient to analyze a single strand of the polynucleotide of interest.

The methods of the current invention can be used to identify the presence of mutations or alterations in the nucleotide sequence of a polypeptide of interest. To identify mutations or alterations, the sequence of the polynucleotide of interest is compared with the sequence of the native or normal polynucleotide. An "alteration" in the polynucleotide of interest, as used herein, refers to a deviation from the expected sequence (the sequence of the native or normal polynucleotide), including deletions, insertions, point mutations, frame-shifts, expanded oligonucleotide repeats, or other changes. The portion of the polynucleotide of interest that contains the alteration is known as the "altered" region. The methods can also be used to determine the polynucleotide sequence of a polypeptide of interest having a previously unknown nucleotide sequence.

In one embodiment of the current invention, the polynucleotide of interest is analyzed by annealing the polynucleotide to an array comprising sets of oligonucleotide primers. The oligonucleotide primers in the array have a length N, where N is from about 7 to about 30 nucleotides, inclusive, and is preferably from 20 to 24 nucleotides, inclusive. Each oligonucleotide primer within each set differs by one base pair. The oligonucleotide primers can be prepared by conventional methods (see Sambrook et al., *Molecular Cloning: A Laboratory Manual* (2nd Ed, 1989)). The sets of oligonucleotide primers are arranged into an array, such that the position and nucleotide content of each oligonucleotide primer on the array is known.

The size and nucleotide content of the oligonucleotide primers in the array depend on the polynucleotide of interest and the region of the polynucleotide of interest for which sequence information is desired. To analyze a polynucleotide of interest for the presence of alterations, consecutive primers differing by one base pair at the growing end and capable of hybridizing successively along the relevant part(s) or the whole of the polynucleotide are used. An example of such a primer set is shown in Figure 1. If only one or a few specific positions of the polynucleotide sequence are examined for alterations, the necessary array of oligonucleotide primers covers only the

mutation regions, and is therefore small. If the whole or a major part of the polynucleotide of interest is to be analyzed for possible mutations at varying positions, the necessary array is larger. For example, the whole hypoxanthine-guanine phosphoribosyl-transferase (HPRT) gene can be covered by 900 primers, arranged in a 30 X 30 array; the whole p53 gene requires 700 primers. If both strands of a double-stranded polynucleotide of interest are analyzed for the presence of alterations, the array comprises consecutive oligonucleotide primers for the suspected mutation region of both the template polynucleotide of interest and the complementary polynucleotide of interest. If the polynucleotide of interest has not been sequenced previously, the array includes oligonucleotide primers comprising all possible N-mers.

The array of sets of oligonucleotide primers is immobilized to a solid support at defined locations (i.e., known positions). The immobilized array is referred to as a "DNA chip", which is the apparatus of the current invention. The solid support can be a plate or chip of glass, silicon, or other material. The solid support can also be coated, such as with gold or silver. Coating may facilitate attachment of the oligonucleotide primers to the surface of the solid support. The oligonucleotide primers can be bound to the solid support by a specific binding pair, such as biotin and avidin or biotin and streptavidin. For example, the primers can be provided with biotin handles in connection with their preparation, and then the biotin-labelled primers can be attached to a streptavidin-coated support. Alternatively, the primers can be bound by a linker arm, such as a covalently bonded hydrocarbon chain, such as a C<sub>10-20</sub> chain. The primers can also be bound directly to the solid support, such as by epoxide/amine coupling chemistry (see Eggers, M.D. et al., Advances in DNA sequencing Technology, SPIE conference proceedings, January 21, 1993). The solid support can be reused, as described in greater detail below.

In another embodiment of the invention, the polynucleotide of interest is analyzed by annealing the polynucleotide to one or more specific oligonucleotide primers that are not attached to a solid support; such oligonucleotide primers are referred to herein as "free oligonucleotide primers". If free oligonucleotide primers are used, the polynucleotide of interest can be attached to a solid support, such as magnetic beads. The free oligonucleotide primers have a length N, as described above, and are prepared by conventional methods (see Sambrook et al., *Molecular Cloning: A Laboratory Manual* (2nd Ed, 1989)). The size and nucleotide content of the free oligonucleotide primers depend on the polynucleotide of interest and the region of the polynucleotide of interest for which sequence information is desired. To analyze a

polynucleotide of interest for the presence of alterations, primers capable of hybridizing immediately adjacent to the relevant part(s) of the polynucleotide are used. If more than one position of the polynucleotide sequence is examined for alterations, the free oligonucleotide primers are mutually distinguishable: i.e., the oligonucleotide primers have different mobilities during gel electrophoresis. In a preferred embodiment, oligonucleotides of different lengths are used. For example, an oligonucleotide primer of 10 nucleotides in length is designed to hybridize immediately adjacent to one putative mutation, and an oligonucleotide primer of 12 nucleotides in length is designed to hybridize immediately adjacent to a second putative mutation. Because the oligonucleotide primers are of different lengths, they will migrate to different positions on the gel. Thus, in this manner, the nucleotide content of each oligonucleotide primer can be identified by the position of the oligonucleotide primer on the gel.

The polynucleotide of interest is hybridized to the array of oligonucleotide primers, or to the free nucleotide primers, under high stringency conditions, so that an exact match between the polynucleotide of interest and the oligonucleotide primers is obtained, without any base-pair mismatches (see Sambrook et al., *Molecular Cloning: A Laboratory Manual* (2nd Ed, 1989)). For example, a schematic illustration of a hypothetical polynucleotide of interest annealed to an oligonucleotide primer that is attached to a solid support is shown schematically in Figure 2. In Figure 2, a part of the sequence of the polynucleotide of interest that follows immediately after the portion of the polynucleotide that is bound to the oligonucleotide primer on the array is shown as TGCAACTA. Six corresponding consecutive primers are shown in Figure 3, i.e. primers ending with the pairing bases A, AC, ACG, etc. If the polynucleotide of interest is double-stranded, it can be separated into two single strands either before or after the binding of the polynucleotide of interest to the array oligonucleotide primers. Both the template and the complementary polynucleotide of interest can be analyzed utilizing a single array. Thus, while not shown in Figure 2, appropriate primers corresponding to the complementary polynucleotide of interest are also attached to the solid support in known positions.

When the polynucleotide of interest is hybridized to the array of sets of oligonucleotide primers, or to the free oligonucleotide primers, under hybridization conditions, annealed primers are formed. The term, "annealed primer", as used herein, refers to an oligonucleotide primer (either free or attached to a solid support) to which a polynucleotide of interest is hybridized. The annealed primers are subjected to a single base extension reaction. The "single base extension reaction", as used herein, refers to a



The terminating nucleotides preferably comprise dNTPs, and particularly comprise dideoxynucleotides (ddNTPs), but other terminating nucleotides apparent to the skilled person can also be used. If the terminating nucleotides are single nucleotides, then nucleotides corresponding to each of the four bases (A, T, G and C) are utilized in the single base extension reaction. If the terminating nucleotides are dinucleotide units, for example, then nucleotides corresponding to each of the sixteen possible dinucleotides are utilized.

The nucleotides are mutually distinguishable. For example, if the solid support is coated with a free electron metal, such as with gold or silver, surface plasmon resonance (SPR) microscopy allows identification of each nucleotide, by the change of the refractive index at the surface caused by each base extension. Alternatively, at least one of the terminating nucleotides is labelled by standard methods to facilitate detection. Suitable labels include fluorescent dyes, chemiluminescence, and radionuclides. The number of nucleotides that are labelled can be varied. It is sufficient to use three labelled terminating nucleotides, the fourth terminating nucleotide being identified by its "non-label", if single nucleotides are added in the base extension reaction. For example, if one is examining the polynucleotide of interest for the presence of a particular alteration, and not for the complete base sequence in the altered region, three labelled terminating nucleotides are sufficient. Fewer than three labels can also be

utilized under appropriate circumstances. An exemplification of the use of two labelled and two unlabelled dNTPs is described below. If a specific alteration is to be investigated, such as a point mutation, only the native or normal nucleotide need be labelled, as a mutation would be indicated by the presence of the "non-label".

- 5 Alternatively, the expected mutant nucleotide can also be labelled.

After the single base extension reaction has been performed, the identity and location of each terminating nucleotide is observed. If free oligonucleotide primers are used, the extended primers are eluted and separated by gel electrophoresis, and the gel is then analyzed. If oligonucleotide primers attached to an array are used, the array itself is  
10 analyzed. The gel or array is analyzed by detecting the labelled, terminating nucleotides bound to the oligonucleotide primers. The labeled, terminating nucleotides are detected by conventional methods, such as by an optical system. For example, a laser excitation source can be used in conjunction with a filter set to isolate the fluorescence emission of a particular type of terminating nucleotide. Either a photomultiplier tube, a charged-  
15 coupled device (CCD), or another suitable fluorescence detection method can be used to detect the emitted light from fluorescent terminating nucleotides.

The sequence of the polynucleotide of interest can be analyzed from the label pattern observed on the array or on the gel, since the position of each different primer on the array or on the gel is known, and since the identity of each terminating nucleotide  
20 can be determined by its specific label. The label and position of each terminating nucleotide either within the array or on the gel will directly define the sequence of the polynucleotide of interest that is being analyzed. Mutations or alterations in the sequence of the polynucleotide of interest are indicated by alterations in the expected label pattern. For example, assume that the nucleotide sequence shown in Figure 2  
25 contains a mutation: the third base C from the left is replaced by a G in the polynucleotide of interest. The top primer in Figure 3 will still be extended by a C as shown in Figure 4, whereas the next primer will be extended by a C rather than a G. Since this new, unexpected base C can be identified by its specific label and the respective primer location is known, the corresponding base mutation is identified as G.

30 The following simple example illustrates the ability to obtain complete sequence information and to identify a mutation in a representative polynucleotide of interest. The example utilizes two labelled terminating nucleotides, which give complete sequence information.

Assume a normal polynucleotide of the following base pair composition:

+ A C T G C T T A G  
- T G A C G A A T C

and a corresponding mutant polynucleotide having the following base pair composition, which has a single base mutation in the third base pair:

5 + A C C G C T T A G  
- T G G C G A A T C.

Using fluorescent labelling, for example, with a red label ("R") for terminating A and a green label ("G") for terminating G, and no label, i.e., null ("N") for the remaining bases T and C, the following "binary" codes allowing sequence interpretation would be  
10 obtained for the normal, mutant and heterozygote sequences, respectively:

	+ N-G-R-N-G-R-R-N-N	Normal
	- R-N-N-G-N-N-N-R-G	
	+ N-G-G-N-G-R-R-N-N	Mutant (Affected)
	- R-N-N-G-N-N-N-R-G	
15	R	
	+ N-G-G-N-G-R-R-N-N	Heterozygote (Carrier)
	- R-N-N-G-N-N-N-R-G.	

The presence of such a point mutation will affect the base pairing of the next few oligonucleotide primers to the polynucleotide of interest, and thereby the primer  
20 extensions obtained, such that the bases in the vicinity of the mutation (i.e., in the altered region) may not be accurately identified. To optimize identification of bases in the altered region, it is preferred to analyze both strands of such a double-stranded polynucleotide of interest. The few bases that may be difficult to identify on the template polynucleotide of interest, as well as the changed base, will be identified by the  
25 base extensions of the primers for the complementary polynucleotide of interest, as the analysis of the complementary polynucleotide of interest approaches the mutation site from the opposite direction. In the nearest regions on either side of the alteration, the sequence determination is thereby provided by the oligonucleotide primers for one of the two strands.

30 The sequence of a polynucleotide of interest for which the sequence is previously known can be determined using methods similar to those described above in reference to identification of mutations utilizing an array of oligonucleotide primers. As before, the positions of the terminating nucleotides within the array will directly define the sequence position of each nucleotide in the polynucleotide of interest.

002447611300

After analysis of the nucleotide sequence of a polypeptide of interest, the polynucleotide of interest and the terminating nucleotides can be removed from the DNA chip, so that the chip can be reused. In a preferred embodiment, the added terminating nucleotides are capable of being removed from the solid support after analysis of the polynucleotide of interest has been completed. Once the nucleotides are removed, the solid support with the immobilized oligonucleotide primers can be used for a new analysis. The nucleotides can be removed using standard methods, such as enzymatic cleavage or chemical degradation. Enzymatic cleavage, for example, would use a terminating nucleotide which can be removed by an enzyme. The single base extension reaction could result in addition to the oligonucleotide primers of RNA dideoxyTTP or RNA dideoxyCTP by reverse transcriptase or other polymerase. A C/T cleavage enzyme, such as RNase A, can then be used to "strip" off the RNA dideoxynucleotides. Alternatively, sulfur-containing dideoxy-A or dideoxy-G can be used during the single extension reaction; a sulfur-specific esterase, which does not cleave phosphates can then be used to cleave off the dideoxynucleotides. For chemical degradation, a chemically degradable terminating nucleotide can be used. For example, a modified ribonucleotide having its 2'- and 3'-hydroxyl groups esterified, such as by

35 a modified ribonucleotide having its 2'- and 3'-hydroxyl groups esterified, such as by

acetyl groups, can be used. After binding of the terminating nucleotide to the annealed primer, the acetyl groups are removed by treatment with a base to expose the 2'- and 3'-hydroxyl groups. The ribose residue can then be degraded by periodate oxidation, and the residual phosphate group removed from the annealed primer by treatment with a  
5 base and alkaline phosphatase.

The method and apparatus of the current invention have uses in detecting mutations, deletions, expanded oligonucleotide repeats, and other genetic abnormalities. For example, the current invention can be used to identify frame shifting mutations caused by insertions or deletions. Furthermore, carrier status of heritable diseases, such  
10 as cystic fibrosis,  $\beta$ -thalassemia,  $\alpha$ -1, Gaucher's disease, Tay Sach's disease, or Lesch-Nyham syndrome, can be easily determined using the current invention, because both the normal and the altered signals would be detected. Furthermore, mixtures of DNA molecules such as occur in HIV infected patients with drug resistance can be determined. The HIV virus may develop resistance against drugs like AZT by point  
15 mutations in the nucleotide sequence of a reverse transcriptase (RT) gene. When mutated viruses start to appear in the virus population, both the mutated gene and the normal (wild type) gene can be detected. The greater the proportion is of the mutant, the greater is the signal from the corresponding mutant terminating nucleotide.

The current invention is further exemplified by the following Examples.

20 EXAMPLE 1 Analyzing the Sequence of a Polynucleotide of Interest Utilizing Free Oligonucleotide Primers

An analysis of the hypoxanthine-guanine phosphoribosyl-transferase (HPRT) gene (the polypeptide of interest) was conducted for three individuals (Patients A, B, and C).

25 A. Obtaining the Polynucleotide of Interest

The polymerase chain reaction (see Sambrook et al., *Molecular Cloning: A Laboratory Manual* (2nd Ed, 1989), especially chapter 14) was utilized to amplify the polynucleotide of interest. During the reaction, one of the two PCR primers was tagged with a biotin group. Following amplification, the single strand template was captured  
30 with streptavidin coated magnetic beads. For a 50  $\mu$ l PCR reaction, 25  $\mu$ l of Dynal M-280 paramagnetic beads (Dynal A/S, Oslo, Norway) was used. The supernatant of the beads was removed and replaced with 50  $\mu$ l of a binding and washing buffer (10 mM

5 150  $\mu$ l of  
0.15 M NaOH was again added for five minutes. Following denaturation, the beads  
were washed once with 150  $\mu$ l of 0.15 M NaOH and twice with 1X T7 annealing buffer  
(40 mM Tris-HCl (pH 7.5); 20 mM  $MgCl_2$ ; 50 mM NaCl). The beads were finally  
suspended in 70  $\mu$ l of water. This process both isolates the single-stranded  
10 polynucleotide of interest and removes any unincorporated dNTPs remaining after PCR.

After single strand isolation, the oligonucleotide primers were annealed to the polynucleotide of interest by heating to 65°C for approximately two minutes and cooling to room temperature over approximately 20 minutes. The 10 µl reaction volume consisted of 7 µl of the polynucleotide of interest (0.5-1 pmol), 2 µl of 5X T7 annealing buffer, and 1 µl of extension primer (3-9 pmol). The extension reaction was then performed. For the reaction, 1 µl of DTT, 2 µl of T7 polymerase (diluted 1:8) and 1 µl of ddNTPs (final concentration of 0.5 uM) were added. The reaction proceeded at 37°C for two minutes, and then was stopped by the addition of 100 µl of washing buffer (1XSSPE, 0.1% SDS, 30% ethanol). The beads were washed twice with 150 µl of the washing buffer. The extension products were eluted by the addition of 5 µl of formamide and heated to 70°C for two minutes. The beads were captured by the magnet and the supernatant containing the extension products was collected and analyzed on a ABI 373 (Applied Biosystems, Inc.). Oligonucleotide primers of lengths varying from 10 to 17 were used. As shown in Figure 5, extension products were formed efficiently.

Each ddNTP was labelled by a different fluorophore. ABI Dye Terminator dyes designed for taq polymerase were used: ddG is blue, ddA is green, ddT is yellow, and ddC is red. Four fluorescent ddNTPs were added to each reaction tube. The extension products were purified, gel separated, and analyzed on an ABI 373. Two different bases of exon 3 of the HPRT gene were analyzed: base 16534 (wild type is A) and base 16620 (wild type is C).

The results of the four fluor, single lane, indicated that the presence of mutations could be identified easily. All three patients are wild type for A at base 16534 (data not shown). Electrophoretograms shown in Figures 6A, 6B and 6C indicate that Patient A is wild type (C) at base 16620 (Figure 6A), patient B is a mutated individual (C-->T) at base 16620 (Figure 6B), and patient C is a carrier at base 16620 (both C and T) (Figure 6C).

## 2. Deoxynucleotide Labelling - Single Fluorophore

Each ddNTP was labelled by the same fluorophore. DuPont NEN fluorescein dyes (NEL 400-404) were used. Each ddNTP appears blue in the ABI 373. Only one fluorescent ddNTP is added to each reaction tube. The extension products were purified, gel separated, and analyzed on an ABI 373. Four lanes on the gel must be used to analyze each base. Two different bases of exon 3 of the HPRT gene were analyzed: base 16534 (wild type is A) and base 16620 (wild type is C).

The results of the single fluor, four lane, demonstrated results that were identical to those obtained using the four fluor, single lane method described in (1), above. This type of assay minimizes the effect of the fluorophore differences during extension product formation and gel separation.

## 3. Deoxynucleotide Labelling - Biotinlyated Dideoxynucleotides

The ddNTPs are labelled with a biotin group. Four separate reactions are performed, whereby only one of the four ddNTPs is biotinylated. Following the extension reaction, a streptavidin (or avidin) coupled fluorescent group is attached to the biotinylated ddNTPs. Because the biotin group is small, uniform incorporation of the ddNTPs is expected and base-specific differences in extension are minimized. Furthermore, the fluorescent signal can be amplified because the biotin group can bind a streptavidin moiety coupled to multiple fluors.

### EXAMPLE 2 Analyzing the Sequence of a Polynucleotide of Interest Utilizing Labelled Deoxynucleotides

An analysis of the hypoxanthine-guanine phosphoribosyl-transferase (HPRT) gene (the nucleotide sequence of a polypeptide of interest) was conducted for three individuals (Patients A, B, and C). The third exon of the HPRT gene was examined.

Microscope glass slides were epoxysilanated at 80°C for eight hours using 25% 3' glycidoxy propyltriethoxysilane (Aldrich Chemical) in dry xylene (Aldrich Chemical)

with a catalytic amount of diisopropylethylamine (Aldrich Chemical), according to Southern (Nucl. Acids Res. 20:1679 (1992), and Genomics 13:1008 (1992)). The DNA chips were made by placing 0.5  $\mu$ l drops of 5'-amino-linked oligonucleotides (50  $\mu$ M, 0.1 M NaOH) at 37°C for six hours in a humid environment. The chips were washed in 50°C water for 15 minutes, dried and used. The annealing reaction consisted of adding 2.2  $\mu$ l of single-stranded DNA (0.1  $\mu$ M in T7 reaction buffer) to each grid position, heating the chip in a humid environment to 70°C and then cooling slowly to room temperature. A 1  $\mu$ l drop of 0.1 M DTT, 3 units of Sequenase Version 2.0 (USB), 5  $\mu$ Ci  $\alpha$ -<sup>32</sup>P dNTP (3000 Ci/mmol) (DuPont NEN) and noncompeting unlabeled 18.5  $\mu$ M ddNTPs (Pharmacia) were added to each grid position for three minutes. The reaction was stopped by washing in 75°C water, and analyzed on a PhosphorImager (Molecular Dynamics).

Figures 7A, 7B and 7C depict the results of a DNA chip-based analysis for a five-base region within the third exon of the HPRT gene. The rows correspond to a particular base under investigation, and the columns correspond to the labeled base. Figure 7A demonstrates the wild type sequence (TCGAG), Figure 7B demonstrates a C->T mutation, and Figure 7C demonstrates a C-->T mutation.

Those skilled in the art will recognize, or be able to ascertain using no more than routine experimentation, many equivalents to the specific embodiments of the invention described specifically herein. Such equivalents are intended to be encompassed in the scope of the following claims.